

Surface Analysis of Alloy 22 Coupons Exposed for five years to Concentrated Ground Waters

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October 17, 2003

CORROSION/2004 New Orleans, LA, United States March 28, 2004 through April 1, 2004

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SURFACEANALYSISOFALLOY22COUPONSEXPOSEDFORFIVEYEARSTO CONCENTRATEDGROUNDWATERS

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ABSTRACT

Alloy22(N06022)isthecandidatematerialforthecorrosionresistant, outerbarrierofthenuclearwaste container. Twoo fthepotential corrosion degradation modes of the container are uniform corrosion and localized corrosion. At esting programathe Lawrence Livermore National Laboratory is being carried out for Yucca Mountain to determine the susceptibility of Alloy 22 to these two forms of corrosion using long-term immersion tests. Metallic coupons were exposed to several electrolyte solutions simulating concentrated ground water from pH 3 to 10 at 60°C and 90°C. This paper summarizes results on the characteristics of the surface deposits as well as the corrosion rate of 122 coupons of Alloy 22 obtained after more than a five -year exposure. The surface deposits consisted primarily of salt components in the respective solutions. Results showed little general corrosion and the absence of localized (crevice) corrosion.

Keywords: N06022, general corrosion, surface deposits, concentrated ground waters

INTRODUCTION

The proposed engineering barriers that will limit the release of radioactive material in the Yucca Mountainrepositorywillconsistofasealedcontainerandadetacheddripshield. The container will be double walled with an internal barrier of type 316L stainless steel (S31603) and an external barrier of Alloy22 (N06022). The dripshield will be madew ith Titanium Grade 7 (R52400). The internal barrier of the container will serve to shield radiation and also provide mechanical integrity. The primary purpose of the outerwall of the container is to provide protection against corrosion. The presence of the dripshield will guard the containers against water see page and rock fall from the drift walls. Alloy 22 (N06022) was selected for the corrosion resistant barrier of the containers because it is well known commercially for its excellent corrosion behavior in aggressive environments.

1-5 Itisnickel -based (Ni) and its nominal composition (weight percent) is ~57% nickel (Ni), 22% chromium (Cr), 13% molybdenum (Mo), 3% tungsten (W) and 3% iron (Fe). Because of its high Cr content, Alloy 22

remains pass ive in most industrial environments and thus, has an exceptionally low general corrosion rate.

In the absence of the drip shield, waters that contact the waste container are expected to be in the form of a multi-ionic solution. These solutions may form through two different mechanisms: (1) Dripping from the drift wall and concentrating on the container surface or (2) Deliquescence of salts that may accumulateontopofthecontainerduringdryperiods. Inbothcases, the aqueous solution are expected to be concentrated and contain multiple component. The ground waters that are associated with the ^{6,7}Table 1 shows the composition of a saturated YuccaMountainregionhavebeenwellcharacterized. zone water (from a well designated, J -13) from near the repository site. The well water, J neutral and bicarbonate -rich with significant concentrations of sulfate, nitrate, chloride, alkalis and alkaline earths ions. Table 1 also shows the composition of various laboratory -prepared, aqueous, concentrated electrolyte solutions in which testing was performed. These electrolyte solutions range frompH~3to10andaredesignatedassimulatedacidifiedwater(SAW),simulatedconcentratedwater ⁵ reported that after a 2 (SCW) and simulated dilute water (SDW). Farmer et al. Alloy 22 coupons in concentrated aqueous electrolytes from pH 2.8 to 10, at both 60°C and 90°C, the ⁻⁴ mpy). Wong et al. average corrosion rate by mass loss was approximately 20 nm/yr (~8 x 10 determined the corrosion rates after a 5 - year immersion in the same concentrated aqueous electrolytes to behigherforcrevicecouponsthanweightlosscoupons, but still low over all with a maximum corrosion rateofonly23nm/yr.

The purpose of the present work was to cha racterize the surface of Alloy 22 coupons exposed for five years to concentrate dground waters.

EXPERIMENTAL

ThecorrosionrateofAlloy22wasdeterminedusingimmersiontestsaccordingtoASTMG31andG1. ASTM G 31 provides guidelines on laborato ry coupon immersion corrosion testing and ASTM G1 provides guidelines on coupon preparation, cleaning and evaluation. Two types of coupons were used. These were labeled (a) weight loss coupons and (b) crevice coupons. The nominal dimensions were 2 inchx 1/8inch(approximately50mmx25mmx3mm)and2inchx2inchx1/8inch(50mm x50mmx3mm), respectively. For each coupon type, there were two variants, wrought (non -welded) and welded. The coupons were fabricated from Alloy22 plates tock and the chemical compositions for the weight loss and crevice coupons as well as the weld filler metal are shown in Table 2. All weight loss coupons were affixed with a ½ "diameter PTFE or ceramic washer while all crevice coupons were affixed to the coupon holder or rack with a ¾ "diameter PTFE or ceramic crevice former (CF). The purpose of the crevice former was to create an environment that might induce crevice corrosion at the contact interface, or under occluded conditions.

The electrolyte sol utions used for the immersion tests were complex solutions that contained multiple ionic species. The six vessels (26 to 30) that housed the test coupons were filled with approximately 1000 liters of the specific electrolyte solution (Table 1). These ser ies of vessels at Lawrence Livermore National Laboratory is called the Long Term Corrosion Test Facility (LTCTF). It is also important to mention that these vessels (26 - 30) not only contained coupons and U -bend specimens of Alloy 22 but

also of other corr osion resistant alloys such as Alloy 825 (N08825), Alloy C -4 (N06455), Alloy G -3 (N06985), Alloy 625 (N06625), Titanium Grades 7,16 and 12 (R52400, R52402 and R53400).

Each of the simulated solutions used in this study were concentrated variations of We ll J -13 water (Table 1). SAW is approximately 1000 times more concentrated than J -13 and is a cidified to pH~3. SDW and SCW are approximately 10 and 1000 times more concentrated than J -13 water, respectively. BothSDWandSCWhaveapH~10. Weightl ossandcrevicecouponsweretestedat60°Cand90°C. Approximately half of the coupons were exposed to the liquid phase of the solution (complete immersion) and the other half were exposed to the vapor phase (suspended over the liquid surface). The coupons were exposed to the testing environments at their free corrosion potential (E external polarization was not applied to the coupons. The coupons were suspended in the test vessels from a non - metallic rack. Each coupon was electrically isolated from the other coupons. The coupons were mounted horizontally flat in the testracks. The front or labels ide of the coupon faced down in the vessel and the backside of the coupon faced up. The reported test temperature corresponded to the liquid phase temperature. In the testing tanks or vessels (LTCTF), the electrolyte solutions were naturally aerated, i.e. the solutions were not purged but the ingress of air above the solution was not restricted. All the tests were carried out under ambient pressure. Welded and non -welded (wrought) coupons were tested in twelve different conditions (3 electrolytes x 2 temperatures x 2 phases). The exposure time for each coupon was approximately 5 years. The actual testing time for each vessel is shownin Table3alongwiththecouponlabel,vesselnumberandaveragecorrosionrateinnm/yr.Each coupon is designated with 3 letters and 3 characteristic numbers. The letter D represents Alloy 22 (N06022),theletterCrepresentscrevicecoupon,theletterW representsweightlosscoupon,theletterA indicates that the coupon is seamless (non -welded) and the letter B indicates that the coupon does contain a weld seam. A total of 134 Alloy 22 test coupons were studied. All of the coupons were individuallys tudiedunderastereomicroscopeupto 100 times magnification. Most of the coupons (122 of 134) were cleaned of surface deposits to measure the corrosion rate. Twelve welded crevice coupons, representingeachofthedifferenttestconditions, were set as ideforsurfaceanalyses. Energy Dispersive Spectroscopy (EDS) was performed on three coupons (DCB054, DCB114 and DCB180), immersed respectively in SAW, SCW and SDW liquid at 90°C. X -ray Photoelectron Spectroscopy (XPS) was performed on the same SAW and SCW coupons (DCB054 and DCB114). The surface analysis results andbriefdescriptionsofthesurfaceappearancearesummarizedinTable4.

After an approximate five -year exposure to each solution/environmental condition, the coupons were removed from the eirrespective test vessel. All 134 coupons were rinsed with de -ionized(DI)waterand stored in pre -labeled, individual containers. Each coupon was then digitally photographed from the front(labelside)andfromtheback.Allcouponswerethenweighed threetimesatdifferenttimesofthe daytoensurecompleteremovalofmoisture. For example, coupon "X" was weighed in the morning and afternoonofDay1andthenweighedagaininthemorningofDay2.Inallofthetestedconditions,the coupons remo ved from the test vessels were covered with deposits. Therefore, the coupons were cleanedpriortofinalweighingforthecalculationofcorrosionratesbyweight(mass)loss.Anexample of one of the weight -loss coupons is shown in Figure 1, before and aftercleaning. The cleaned surface is deposit-free and illustrates the effectiveness of the cleaning solutions used. Figure 1 also shows that the coupon did not suffer noticeable corrosion in the tested electrolyte. All coupons tested in alkaline solutions (SCW and SDW) were cleaned per ASTMG 1 -C6.1, which specifies cleaning with a HCl solution at ambient temperature. The immersion time for cleaning was approximately 2 min. All couponstestedinanacidifiedsolution(SAW)were cleaned per ASTMG1 -C6.1 and C7.4. That is, a

acid cleaning was followed by an alkaline cleaning. After the coupons were immersed in the HCl solution, they were rinsed in DI water and later immersed in hot (90 $\,$ -95°C) sodium hydroxide and potassium permanganate (NaOH/KMnO $_4$) for 3 min. followed by a cleaning step using a diammonium citrate ((NH $_4$)₂HC $_6$ H $_5$ O $_7$) solution at ambient temperature for 2 min. In an attempt to characterize some of the possible soluble corrosion products, aliquots of the solutions used for cleaning were ana lyzed using ICP -MS (Inductively Coupled Plasma Mass Spectroscopy) and ICP -AES (Inductively Coupled Plasma – Atomic Emission Spectroscopy) techniques. The solution analysis results are also shown in Table 4.

The corrosion rates of the cleaned coupons we rethen calculated using Equation 1,

$$CR(nm/yr) = \frac{8.76x10^{-10} \cdot \Delta W}{\rho \cdot A \cdot t}$$
 (1)

where 8.76×10^{-10} is the proportionality constant, ΔW is the mass loss in grams after 5+ years, pisthe density of Alloy 22 (8.69 g/cm⁻³), A is the exposed surface area of each coupon (cm⁻²) and t is the exposure time (hours).

RESULTSANDDISCUSSION

DepositsontheImmersedCoupons

As stated above, after removal from the test vessels, all coupons were covered with a varied degree of deposits (Table 4). In general, these deposits gave the coupons a characteristic appearance that varied from vessel to vessel. For example, the coupons exposed to the SAW solution were golden/brown on the label side (facing down in the vessel) and bluish/gree nonthe backside (facing up). On the other hand, the coupons exposed to the SCW solution were gray in appearance and had a thick layer of white, salt-like deposits on the backside (facing up) (Table 4). Table 5 shows the amount of accumulated deposits ineachtestedcondition. That is, the data in Table 5 shows the weight of the dry coupons after removal from the vessels minus the weight of the coupons before immersion in the vessels in 1997. In the vapor phase, more deposits formed at 60°C than at 90 °C, and at each testing temperature, more deposits generally formed at the liquid than at the vapor phase. In a few cases, the coupons experienced weight loss even before cleaning. Table 5 shows that coupons immersed in SCW liquid at 90°C exhibited approximately 10 times the weight gain than those coupons immersed in SAW at the same temperature. This is due to the large amount of white deposits in SCW described in Table 4. The white temperature are the contraction of the condepositwasmostlycalciumcarbonate(CaCO

Table 4 briefly descri bes the surface appearance of the coupons from each vessel. The coupons were mostly dull, light blue -gray or tan; however, those immersed in the SAW exhibited blue and iridescent green backsides. This discoloration appears to be dependent on the vesselo rsolution chemistry. Other predominant features observed include rust -like deposits, mostly in the label region, white salt -like deposits scattered throughout the coupon surface and generally, a clearer crevice former (CF) annuli. That is, a lesser amou nto f deposits formed under the crevice former than on the surface exposed to the bulk of the solution. Similarly, the crevice former annulus of each coupon was shinier metallic in

appearance than the surface exposed boldly to the solution. That is, them seemed to have interactedless with the environment than the rest of the coupon.

Figure 2 shows some of the rust -like deposits residing in the label lettering or grooves. EDS analyses confirmhighamountsofironandoxygen (iron -oxideorrust)inthedeposits at the label location. It is likely that this iron was transferred to the coupon by the labeling tool (stamp) or letter indenter. Iron oxides(hydroxides)wereobservedinthelabelregionsincouponsexposedtoall thevessels. However, in the acidic solutions (SAW in vessels 25 and 26), iron seemed to have dissolved more in the electrolyte and later deposited throughout all the coupons in the vessels giving them the characteristic goldencolor(Table 4). On the ot herhand, their on present in the grooves of the label of the coupons exposed to vessels 27 to 30 (alkaline SCW and SDW) was more in soluble, that is, it stayed localized in the grooves and did not spread as much over the entire surface of the coupons in th ese vessels. Therefore, the coupons in the alkaline solution vessels had shades of gray (Table 4). Figure 3 shows SEMimages of some of the other deposits that were observed on the coupons immersed in (a) SAW at 90°C and (b) SCW at 90°C. A summary of th e deposits, as detected by EDS and XPS, is shown in Table 4. There is generally good agreement between the two analytical methods. Table 4 shows that ironispresentontheentiresurfaceofthecouponsexposedtoSAW; however is not present in the bold surface of the coupons exposed to SCW. EDS and XPS also detected the presence of silicon (Si), aluminum (Al) and magnesium (Mg) in the coupons exposed to SCW (Table 4). Si could have co precipitatedfromtheelectrolyte(Table 1); however, the presence o fMgandAlinthecouponsexposed toSCWcannotbefullyexplained. They could have been transferred withir on by the labeling tool. The deposits on the coupons exposed to SCW also contained fluoride, probably as calcium fluoride (CaF 2), which is a rath er insoluble compound. The chemical analysis of aliquots of the solutions used for cleaningshowedthepresenceofcations(suchasCa)thatwerepartofthetestsolution,confirmingthat thedepositsformedbyprecipitationofsaltsfromthetestingele ctrolytes.

Using ICP -MS and ICP -AES techniques, only silicon (Si), calcium (Ca) and magnesium (Mg) were detected in significant or reliable quantities. Si was observed in the NaOH/KMnO 4 solutions used to cleanthecouponstestedinSAW. This was expec tedsinceSiO 2mayhaveprecipitatedonthecoupons in the acidic SAW and later was removed by cleaning in the caustic solution. Table 4 shows that both EDS and XPS analyses detected high concentrations of Si and O in the deposits found in Vessel 26 (SAW,90°C). Itwassurprising to detect a small amount of SiO 2inthediluteHClsolutionusedforthe first cleaning of the coupons tested in 90°C liquid SCW since SiO 2 is not soluble in acidic solutions. Silica(SiO₂)isnotexpectedtoprecipitate from an equilibrated SCW solution. High concentrations of Cawere found in all HCl solutions that were used to clean the coupons tested in SDW, those immersed in 90°C SCW and in some of the HCl and diammonium citrate solutions used to clean the coupons tested in SAW. With the exception of the citrate cleaning, these calcium results were also anticipated since CaCO₃ is expected to precipitate on the coupons from equilibrated SDW and SCW solutions. Table 4 shows that both EDS and XPS analyses detected significant amounts of Ca, C and O in the deposits found in Vessel 28 and 30 (SDW solution). The addition of a cid (e.g. HCl) to CaCO 3ledtothe formationofcalciumionsandtheevolutionofCO 2gas.

CorrosionRates

Table3showsthecalculated, average corrosi on rates for the Alloy22 weight loss and crevice coupons exposed to the SAW, SCW and SDW solutions at 60°C and 90°C for over 5 years. The average

corrosionrates and standard deviations are presented. Even though the optical appearance and amount of deposits on coupons exposed to different solutions were different, the calculated corrosion rate was similar for coupons exposed to different vessels. The lowest rates for the weight loss coupons were observed in the SDW solution (Table 3). The highest rate sobserved in the SDW solution vessels (Table 3). In most cases, the crevice coupons exhibited corrosion rates 2 -5 timeshigher than the weight loss coupons in the same solutions. Stereomicroscopic and scanning electron microscope (SEM) observations of both weight loss and crevice coupons indicated little or no corrosion for Alloy 22. After 5 years immersion, the machining grooves remained uniform and sharp throughout each coupon (Figures 1 and 4).

Sincepreferential dissolution was not observed below the crevice former annuli of the Alloy 22 crevice coupons, it was previously not clearly understood why the overall corrosion rates of the crevice coupons were generally higher than the corrosion rates of the weight loss coupons. One possible explanation is shown in Figure 4, which shows the front side and backside surfaces of the crevice coupon DCB 081. The front side and backside of the crevice coupons were especified to have a surface finish of RMS 16 (approximately 240 grit), however the backside of the crevice coupons had a surface finish that is not characteristic of paper grinding. This unusual aspect of the surface may have contributed to the difference in corrosion rates between the weight loss and crevice coupons. By comparison, both sides of the weight loss coupons exhibit a surface roughness of RMS 32 (approximately 150 grit) (Figure 1).

Figure 5 shows a schematic representation of the welded weight loss and crevice coupons. Half of the tested coupons contained a GMAW (Gas Metal Arc Weld) seam. After 5 years of immersion testing, none of the welded coupons showed any indication of preferential weld corrosion or weldetching. The weld location was never discernible in the coupons either by optical stereous relectron microscopy. It has also been previously reported that the welded coupons did not shown hanced general corrosion rate over the non-welded coupons.

Finally, in all coupons, the crevice former annuli were clear and absent of any crevice corrosion. The higher corrosion rates observed for the crevice coupons are attributed to the difference in surface finish between weight loss (Figure 1) and crevice (Figure 4) coupons and not to crevice corrosion.

ACKNOWLEDGEMENTS

This work was performed under the auspices of the U. S. Department of Energy by the University of California Lawrence Livermore National Laboratory under contract No. W -7405-Eng-48. This work is supported by the Yucc a Mountain Project -LLNL, which is part of the Office of Civilian Radioactive WasteManagement (OCRWM).

CONCLUSIONS

- (1) After removal from the tanks, the color of the Alloy 22 coupons from each vessel varied as a consequence of the different types of deposits formed on them. For example, salt—like white and golden crystals containing iron.
- (2) EDS, XPS, ICP -MS and ICP -AES techniques showed that the surface deposits were generally compounds that can be traced to the original solution components.
- (3) Alloy2 2couponsexposedtoSAW,SCWandSDWsolutionsat60°Cand90°Cforover5years exhibitedverylowcorrosionrates.Themaximummeasuredcorrosionratewas23nm/yr.
- (4) Thecrevicecouponsgenerallyexhibitedcorrosionrates 2 -5timeshigherthanthesm allerweight loss coupons. Stereomicroscopic observations and scanning electron microscopy indicated little or no general corrosion and the absence of crevice corrosion in all the coupons. The higher corrosionrates observed for the crevice coupons appear to be due to the different surface finish and not to crevice corrosion.

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 $TABLE1\\ CHEMICAL COMPOSITION OF THE ELECTROLYTES OLUTIONS (mg/L)$

Ion	SDW	SCW	SAW	J-13WellW ater
	pH10.1	pH10.3	pH2.8	pH7.4
K ⁺	34	3400	3400	5.04
Na ⁺	409	40,900	40,900	45.8
Mg^{2+}	1	<1	1000	2.01
$egin{aligned} \mathbf{Na}^+ \ \mathbf{Mg}^{2+} \ \mathbf{Ca}^{2+} \end{aligned}$	0.5	<1	1000	13
F	14	1400	0	2.18
Cl	67	6700	24,250	7.14
NO_3	64	6400	23,000	8.78
NO_3 SO_4^{2-}	167	16,700	38,600	18.4
HCO_3^-	947	70,000	0	128.9
SiO ₂ (aq)	~40	~40	~40	61.1

 $TABLE2\\ CHEMICAL COMPOSITION OF THE STUDIED ALLOY 22 HEATS (Wt\%)$

Element	WeightLoss/	WeightLoss/	WeldFiller	WeldFiller
	Crevice	Crevice	Metal	Metal
	(Heat 2277-0-3264)	(Heat2277 -5-3203)	(Heat2277 -4-3263)	(Heat2277 -4-3142)
С	0.004	0.002	0.002	0.003
Co	1.14	1.82	0.89	2.03
Cr	21.3	21.3	21.6	21.17
Fe	4.4	4.0	3.6	4.24
Mn	0.29	0.19	0.32	0.25
Mo	13.4	13.08	13.5	13.81
Ni	~56	~56	~56	~55
P	0.01	0.005	0.009	0.009
S	< 0.002	0.008	0.003	0.005
V	0.17	0.14	0.15	0.13
W	2.9	2.93	2.9	3.01

TABLE3:LISTOFEX AMINEDCOUPONSANDA VERAGECORROSIONRAT ES(nm/yr)

	SAW,	SAW,	SCW,	SCW,	SDW,	SDW,
	60°C	90°C	60°C	90°C	60°C	90°C
Vessel	25	26	27	28	29	30
Datein	06Feb1997	21Feb1997	10Mar1997	10Apr1997	14Apr1997	05Jun1997
Dateout	20May2002	21May2002	17May2002	22May2002	10May2002	22May2002
Exp.Time,days	1930	1916	1895	1869	1853	1813
(h)	(46,320h)	(45,984h)	(45,480 h)	(44,856h)	(44,472h)	43,512h)
WeightLoss -	DWA019	DWA059	DWA089	DWA129	DWA147	DWA174
VaporPhase	DWA020	DWA060	DWA090	DWA130		
1	DWA021	DWA061	DWA091	DWA131		
Welded	DWB019	DWB059	DWB089	DWB129	DWB147	DWB174
WeightLoss -	DWB020	DWB060	DWB090	DWB130		
VaporPhase	DWB021	DWB061	DWB091	DWB131		
	40.40					
AvgRate±s	1.9 ±1.8	1.5 ±1.2	0.4 ±1.2	2.1 ±1.0	0.4 ±0.5	1.5 ±1.1
WeightLoss -	DWA022	DWA062	DWA092	DWA132	DWA148	DWA175
LiquidPhase	DWA023	DWA063	DWA093	DWA133		
	DWA024	DWA064	DWA094	DWA134		
Welded	DWB022	DWB062	DWB092	DWB132	DWB148	DWB175
WeightLoss -	DWB022 DWB023	DWB063	DWB093	DWB132 DWB133	DWD140	DWD173
LiquidPhase	DWB023 DWB024	DWB064	DWB094	DWB133		
Elquidi nuse	D W B024	D W Boot	DWBOJT	DWB134		
AvgRate±s	2.8±1.4	2.3±1.3	3.2±1.9	9.5±2.4	1.1±0.5	0.8±1 .1
WeightLoss -	DWA034	DWA039	DWA104	DWA109	DWA154	DWA167
Waterline	3.6	4.3	2.2	0.0	3.0	2.3
Crevice –	DCA019	DCA049	DCA079	DCA109	DCA139	DCA175
VaporPhase	DCA020	DCA050	DCA080	DCA110	DCA140	DCA176
	DCA021	DCA051	DCA081	DCA111	DCA141	DCA177*
Welded	DCB019	DCB049	DCB079	DCB109	DCB139	DCB175
Crevice -	DCB020	DCB050	DCB080	DCB110	DCB140	DCB176
VaporPhase	DCB021(NA)	DCB051(NA)	DCB081(NA)	DCB111(NA)	DCB141(NA)	DCB177(NA)
			,			
AvgRate±s	8.7 ±1.3	15.1 ±2.0	4.0 ±2.3	5.9 ±3.5	2.8 ±1.7	1.0 ± 0.7
Crevice -	DCA022	DCA052	DCA082	DCA112	DCA142	DCA178
LiquidPhase	DCA023	DCA053	DCA083	DCA113	DCA143	DCA179
	DCA024	DCA054	DCA084	DCA114	DCA144	DCA180
Welded	DCB022	DCB052	DCB082	DCB112	DCB142	DCB178
Crevice -	DCB022 DCB023	DCB052 DCB053	DCB082 DCB083	DCB112 DCB113	DCB142	DCB176
LiquidPhase	DCB024(NA)	DCB054(NA)	DCB084(NA)	DCB114(NA)	DCB144(NA)	DCB180(NA)
AvgRate±s	10.3 ± 7.0	6.1 ±1.5	12.9 ±4.8	8.9 ±3.3	5.6 ± 2.3	4.1 ±3.3

^{*} Data(outlier)isnotincludedintheaverageandstandarddev iation.Thusfar,astatisticaltreatmentusingtheEmpirical CumulativeDistributionFunction(ECDF)appearstoshownosignificanteffectontheresultswhentheoutlierisremoved.

NA = Not available for corrosion rates. (Reserved f s = Standard Deviation

orsurfaceanalyses.)

TABLE4 STEREOMICROSCOPEOBS ERVATIONSOFTHECOU PONS, 2x2COUPONS(WITHACREVICEFORMERATITSCENTER)

Conditions	VaporPhase	LiquidPhase	Surface Deposits (EDS*)	Surface Deposits (XPS ⁺)	Solution Products (ICP-MSand ICP-AES [£])
Vessel25 SAW,60°C	Shiny,bluish -lightgray withbrown/rustdeposits. Rustprevalentinlabeland edges.Fewwhitedeposits aroundCF *.Nocrevice corrosion.	Gray,blue -greenandtanwithrings ofwhiteandbrowndepositsaround hole.Crevicecoupo nswere golden/lightbrownwithaclear, shinyCF.Fewwhite,salt -like depositsunderCF.Bluebackside withdullCF.Nocrevicecorrosion.	NA	NA	Si,Ca
Vessel26 SAW,90°C	Shiny,gray,spottedand highlystained.Isolated browndepositsand/or veil oftanandwhitedeposits. DullanddarkgrayCF.No crevicecorrosion.	Dark,goldenbrownwithbrown deposits.Lightiridescentgreen - goldenbackside.GrayCFwithred edging.Abundantrust/brown deposits.Nocrevicecorrosion.	C,O,S,K,Si Cr,Mo,Fe,W, Ni and<1wt.%: Na,P,Al,Cl,V, Mg,Mn,Co	C,O,Al,Si, Cr,Fe and<1wt.%: Na,P,F,S, Mo	Si,Ca
Vessel27 SCW,60°C	Bluish-purpleand/or golden/yellow.Thinveil ofwhiteortandeposits. Highlyspottedcrevice couponsindicatedroplet condensation.Fewwhite deposits.Noweldetching orcrevicecorrosion.	Shiny,lightgraywiththinveilof graydeposits.Somewhitedeposits aroundringarea.Backsideisduller graywithfewblackandwhite depositsunderCF. Noweldetching. Nocrevicecorrosion.	NA	NA	
Vessel28 SCW,90°C	Shiny,bluish -lightgray, tanandyellowspotted. Thinveiloflightgrayor whitedeposits.Saltsand blueringaroundclearCF. Rustdepositsonlabel.No crevicecorrosion.	Dulldarkgray.Thinlayerofwhite depositsonbothsides(more concentratedonbackside).Shiny, clearCFonlabelsideandlight brownCFonbackside.Nocrevice corrosion.	C,O,Si,F,Mg, Ca,Al,Ni and<1wt.%: Na,Ti,Cl,P, Cr,Mo,Fe,W,	C,O,Si,F, Mg,Ca,Al And<1 wt.%: Na,Ti,Zn& Ni	Si,Ca,Mg
Vessel29 SDW,60°C	Spottedbluish -grayandtan withveilofwhitedeposits. Somerustinlabel.Shiny, lightgrayCF.Dull,dark gray,tanandbluish backsidewithdullCF.Few whitedeposits(thinveil). Nocrevicecorrosion.	Shiny,lighttanorgraywithtanand bluepatches.Smallveilsofwhite deposits.Somerustinlabel.Shiny, lightgrayCFonfrontsideanddull, darkgrayCFonbacksideofcrevice coupons.Nocrevicecor rosion.	NA	NA	Ca
Vessel30 SDW,90°C	Dullwithlightanddark gray,tanandbluishspots andpatches. Veilofwhite deposits. Largewhite depositsandtransparent crystalsaroundhole. Some rustinlabel. Bluish/Gray CFwithrainbowring. No crevicecorrosion.	Dull,darkgraylabelsidewith abundantwhitedeposits.Shiny, lightgray/tanCF.Lightgray -bluish backside.Dull,darkgrayCFwith rainbowringonbackside.No crevicecorrosion.	C,O,Na,Si, Mg,Ca,Cr,W, Ni and<1wt.%: Al,S,P,Cl,Ti, Mo,Fe,Co	NA	Ca

^{*}CF=CreviceFormer(annulus)

[#]EDS=EnergyDispersiveSpectroscopy

⁺XPS=X -rayPhotoelectronSpectroscopy £ICP -MS=InductivelyCoupledPlasma -MassSpectroscopy;ICP -AES=InductivelyCoupledPlasma -AtomicEmissionSpectroscopy

TABLE5
AVERAGEAMOUNTOFDE POSITS(WEIGHTGAIN) ONTHETESTED
ALLOY22COUPONS(mg)BEFORECLEANING

	SAW, 60°C	SAW, 90°C	SCW, 60°C	SCW, 90°C	SDW, 60°C	SDW, 90°C
Vessel	25	26	27	28	29	30
WeightLoss Coupon – VaporPhas e Welded& Nonwelded AvgWtGain±s	1.1±0.5	0.7±0.4	1.5±0.5	0.7±0.4	NA	10.8±4.5
WeightLoss Coupon – LiquidPhase Welded& Nonwelded AvgWtGain±s	8.6±3.0	5.4±0.5	1.5±1.3	44.7±12.4	NA	10.3± 0.6
WeightLoss - Waterline	3.5	2.2	3.7	3.5	NA	53.7
CreviceCoupon - VaporPhase Welded& Nonwelded AvgWtGain±s	1.2±0.7	-1.8±0.3	0.7±0.4	0.7±0.6	1.0±0.3	4.6±5.0
CreviceCoupon - LiquidPhase Welded& Nonwelded AvgWtGain±s	5.8±0.5	8.2±0.9	-0.4±0.8	120.6±5.6 *	0.2±0.4	25.5±0.7

 $^{{\}color{blue}*} \quad \quad \text{Data} (outlier) is not included in the average and standard deviation. There is an apparent measurement error.$

NA=Notavailable.

s = Standard Deviation

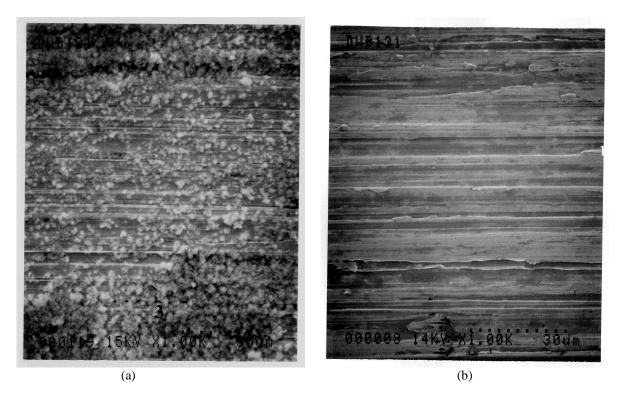
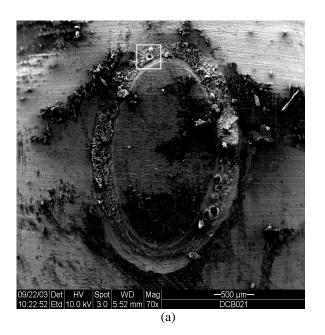
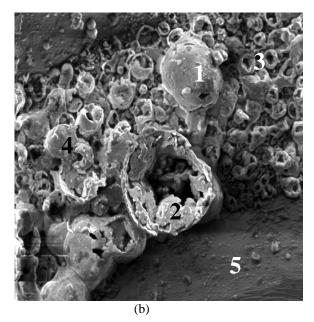


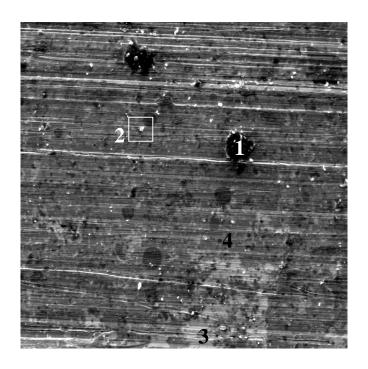
FIGURE1. Alloy22weight -losscoupon,DWB131(90°C,SCW,Vapor), (a)beforeand(b)aftercleaning(1000Xmagnification).

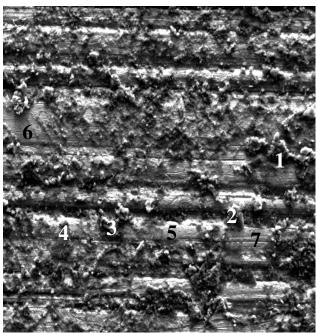




Element	Deposit1 Wt.% (1)	Deposit2 Wt.% (2)	Deposit3 Wt.% (3)	Deposit4 Wt.% (4)	Base1 Wt.% (5)	Alloy22 (nominal) Wt.%
С	19.31	26.89	25.91	21.23	24.88	
О	33.43	35.92	32.47	32.70	11.35	
Si	0.47	0.34	0.53	0.41		
Al			0.28			
Cl	0.27	0.73	0.29	0.15		
S	0.16	1.06	0.75	0.60	1.19	
P		0.22	0.27			
Cr	0.77	0.53	1.07	0.90	12.79	22
Mo	2.48	1.99	2.76	1.05	7.04	13
Fe	40.24	30.76	33.94	40.79	9.01	3
W					3.37	3
Co	2.88	1.56	1.72	2.18	1.04	2
Ni					29.32	57

 $Figure 2. \quad EDS analyses \quad of rust \ -like deposits in stamped label of DCB 021 (90 ^{\circ}C, SAW, Vapor) \\ \quad (a) 70 X magnification (b) 1000 X magnification.$





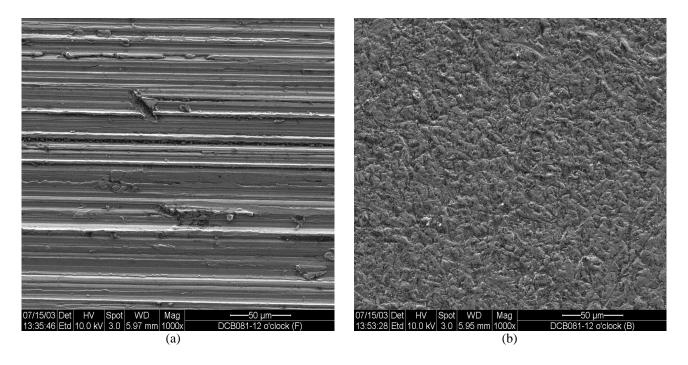
 $\textbf{(a)} DCB054Bold \qquad \textbf{-90}^{\circ}C, Liquid, SAW(pH\textbf{-3})$

Element	Deposit1 Wt.% (1)	Deposit2 Wt.% (2)	Base1 Wt.% (3)	Film Wt.% (4)	Alloy 22 Wt.%
С	58.29	10.28	5.21	9.76	
О	12.82	30.98	4.83	15.86	
Na	0.19	0.80			
Si	0.10	10.61			
Al	0.35	0.93	0.29	0.48	
S	0.61	0.71		1.97	
Mg		0.20			
P	0.36	0.34	0.47	0.53	
Cr	6.11	8.21	19.29	15.59	22
Mo	2.71	5.22	8.76	6.65	13
Fe	9.57	7.91	4.83	10.23	3
W	2.44	9.72	5.53	5.48	3
Co	0.59	0.74	1.60	0.59	2
Ni	5.87	13.34	47.68	32.86	57

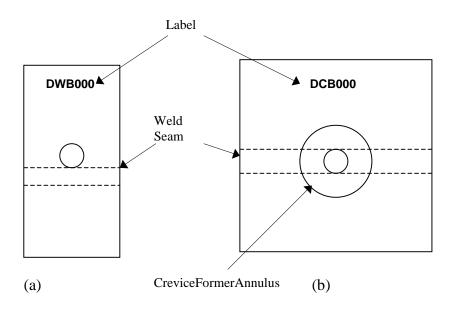
(b)DCB114Bold -90°C,Liquid,SCW(pH~10)

	Deposits	Base	Alloy22
Element	Wt.%	Wt.%	(nominal)
	(1-4)	(5-7)	Wt.%
С	57-81	8-16	
0	12-22	7.7-8.2	
Na	0.3-0.8		
Si	1-5		
Al	0.3-0.8	0.4-0.5	
S		2-3	
Mg	0.5-4.0		
F	0.5-4.0		
Ca	1-4		
Cl	0.1		
S		2-3	
P	0-0.3	0-0.5	
Ti	0.4-4.0		
Cr	0.2-0.9	17-19	22
Mo	0.2-1.1	9-12	13
Fe	0.1-0.3	3.5-4.1	3
W	0-3	6.5-7.4	3
Co	0.1	1.1-1.7	2
Ni	0.4-1.4	35-39	57

Figure 3. EDS analyses of (a) DCB 054 and (b) DCB 114.



 $FIGURE4: SEM images of crevice coupon, DCB081 (60 ^{\circ}C, SCW, Vapor). \\ (a) the front side (b) the back side (1000 X magnification).$



 $FIGURE5 \ -Schematic of Welded (a) Weight Loss and (b) Crevice Coupons.$